Uploading C:\Program Files\Stnexp\Queries\10589864a.str

chain nodes :
6 7 9 15 16 17 19
ring nodes :
1 2 3 4 5 10 11 12 13 14
chain bonds :
2-6 4-7 5-9 11-16 12-17 13-15 14-19
ring bonds :
1-2 1-5 2-3 3-4 4-5 10-11 10-14 11-12 12-13 13-14
exact/norm bonds :
1-2 1-5 2-3 3-4 4-5 5-9 10-11 10-14 11-12 12-13 12-17 13-14 14-19
exact bonds :
2-6 4-7 11-16 13-15

G1:C1,Br

Match level:
1:Atom 2:Atom 3:Atom 4:Atom 5:Atom 6:CLASS 7:CLASS 9:CLASS 10:Atom 11:Atom 12:Atom 13:Atom 14:Atom 15:CLASS 16:CLASS 17:CLASS 19:CLASS fragments assigned product role:
containing 1
fragments assigned reactant/reagent role:
containing 10

L1 STRUCTURE UPLOADED

=> d L1 HAS NO ANSWERS L1 STR

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

Structure attributes must be viewed using STN Express query preparation.

=> s 11 full

FULL SEARCH INITIATED 14:01:28 FILE 'CASREACT'

SCREENING COMPLETE - 98 REACTIONS TO VERIFY FROM 9 DOCUMENTS

100.0% DONE 98 VERIFIED 8 HIT RXNS 1 DOCS

SEARCH TIME: 00.00.01

L2 1 SEA SSS FUL L1 (8 REACTIONS)

=> d 12

L2 ANSWER 1 OF 1 CASREACT COPYRIGHT 2008 ACS on STN

RX(1) OF 20

Cl
$$\stackrel{H}{N}$$
 NO_2 $NaI, Water$ NO_2 $NaI, Water$ NO_2 $NaI, Water$ NaI, Wat

REF: PCT Int. Appl., 2005077913, 25 Aug 2005

NOTE: regioselective CON: 35 hours, 102 deg C

=> d 12 ibib abs

L2 ANSWER 1 OF 1 CASREACT COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 143:229859 CASREACT <<LOGINID::20080920>>

TITLE: Producing 4-nitroimidazole compounds

INVENTOR(S): Shinhama, Koichi

PATENT ASSIGNEE(S): Otsuka Pharmaceutical Co., Ltd., Japan

SOURCE: PCT Int. Appl., 89 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PAT	TENT	NO.		KI	ND .	DATE			A.	PPLI	CATI	и ис	٥.	DATE			
WO 2005077913			A1 20050825				WO 2005-JP2668				 8	20050215					
	W:	ΑE,	AG,	AL,	ΑM,	ΑT,	ΑU,	AZ,	BA,	BB,	BG,	BR,	BW,	BY,	BZ,	CA,	CH,
		CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	ΚE,	KG,	KP,	KR,	KΖ,	LC,	LK,
		LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MΖ,	NA,	ΝI,	NO,
		NΖ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	ТJ,
		TM,	TN,	TR,	TT,	TZ,	UA,	UG,	US,	UΖ,	VC,	VN,	YU,	ZA,	ZM,	ZW	
	RW:	BW,	GH,	GM,	KΕ,	LS,	MW,	MZ,	NΑ,	SD,	SL,	SZ,	TZ,	UG,	ZM,	ZW,	ΑM,
		ΑZ,	BY,	KG,	KΖ,	MD,	RU,	ΤJ,	TM,	AT,	BE,	BG,	CH,	CY,	CZ,	DE,	DK,
		EE,	ES,	FΙ,	FR,	GB,	GR,	HU,	IE,	IS,	ΙT,	LT,	LU,	MC,	NL,	PL,	PT,
		RO,	SE,	SI,	SK,	TR,	BF,	ΒJ,	CF,	CG,	CI,	CM,	GΑ,	GN,	GQ,	GW,	ML,
		MR,	ΝE,	SN,	TD,	ΤG											

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AU	2005212093	B2	20080403				
CA	2555372	A1	20050825	CA 2005-2555372	20050215		
EP	1720838	A1	20061115	EP 2005-710450	20050215		
EP	1720838	B1	20070704				
	R: AT, BE,	BG, CH	, CY, CZ, DE,	DK, EE, ES, FI, FR,	GB, GR, HU, IE,		
	IS, IT,	LI, LT	, LU, MC, NL,	PL, PT, RO, SE, SI,	SK, TR		
CN	1922154	A	20070228	CN 2005-80005310	20050215		
AT	366241	T	20070715	AT 2005-710450	20050215		
BR	2005007777	Α	20070717	BR 2005-7777	20050215		
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IN	2006KN02205	A	20070525	IN 2006-KN2205	20060804		
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US	20070161802	A1	20070712	US 2006-589864	20060817		
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PRIORITY	Y APPLN. INFO	.:		JP 2004-41381	20040218		
				JP 2004-278999	20040927		
				WO 2005-JP2668	20050215		

OTHER SOURCE(S): MARPAT 143:229859

GΙ

AB The present invention provides a method for producing a 4-nitroimidazole (I, X1 = H) at high yield and at high purity by a safe method causing few dangers such as explosion. The production method comprises iodinating a 4-nitroimidazole compound I (wherein each of X1 and X2 represents a Cl or Br), and then reducing the obtained I (X1 = I and X2 is the same as defined above). E.g., 2-bromo-5-iodo-4-nitroimidazole was prepared from 2,5-dibromo-4-nitroimidazole and NaI and the product treated with PtO in the presence of triethylamine in ethanol to give 2-bromo-4-nitroimidazole.

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d 12 1-8

L2 ANSWER 1 OF 1 CASREACT COPYRIGHT 2008 ACS on STN

RX(1) OF 20

REF: PCT Int. Appl., 2005077913, 25 Aug 2005

NOTE: regioselective

CON: 35 hours, 102 deg C